HO5B33/22-HO1B3/12-:

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H05B33/22

European Patent Office Office européen des brevets

(1) Publication number:

145 470

АЗ

EUROPEAN PATENT APPLICATION

(21) Application number: 84308539.0

(51) Int. Cl.4: H 05 B 33/22

(22) Date of filing: 07.12.84

(30) Priority: 09.12.83 JP 233015/83

(43) Date of publication of application: 19.06.85 Bulletin 85/25

(88) Date of deferred publication of search report: 03.06.87

(84) Designated Contracting States: DE FR GB

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(54) Thin-film electroluminescent element.

(57) The development of a dielectric thin-film which is high (140 MV/cm or above) in product of dielectric constant  $\epsilon_i$  and dielectric breakdown field strength E, is essential for realizing an EL element which can operate stably at a low voltage. Such dielectric film is also required which can withstand heat treatments at high temperatures above 500°C and is proof against clouding and in which the electrical breakdown caused by a minute fault produced in the process of film formation is self-healed. A film material which satisfies all of these requirements could be obtained from a TiO<sub>2</sub>-BaO based composition by partially substituting the position of Ti with Sn, Zr of Hf and also partially substituting the position of Ba with Ca or Mg. By using these dielectric films, it is possible to obtain a low-voltage drive thin-film electroluminescent element which are high in production yield and reliability.

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## THIN-FILM ELECTROLUMINESCENT ELEMENT

## 1 TECHNICAL FIELD

The present invention relates to an electroluminescent element, and more particularly to a thin-film
electroluminescent element which is actuated in an AC

5 field. Such electroluminescent element has specific
characteristics that enable the realization of plate
displays, and it is especially suited for adaptation to
character and graphic terminal displays for personal
computers, etc., therefor can be widely applied to the

10 field of office automation systems.

#### BACKGROUND ART

Generally, electroluminescent element (hereinafter abbreviated as EL element) which emitts light upon
application of an AC field has a structure in which a

15 filmy layer of a dielectric is provided on one side or
both sides of a thin layer of an electroluminescent phosphor and these laminate layers are sandwiched by two
electrode layers. The phorphor layer used in such element
is basically composed of such material as ZnS, ZnSe or

20 ZnF<sub>2</sub> doped Mn or a rare-earth fluoride as luminescent
center in said base material. ZnS phosphor element using
Mn as luminescent center is capable of providing a
luminance of up to about 3,500-5,000 Cd/m<sup>2</sup> by the application of an AC voltage with a frequency of 5 kHz.

- Typical examples of dielectric material used in said element are  $Y_2O_3$ ,  $SiO_2$ ,  $Si_3N_4$ ,  $Al_2O_3$  and  $Ta_2O_5$ . The thickness of ZnS layer is about 5,000 to 7,000 Å and that of dielectric layer is about 4,000 to 8,000 Å.
- In the case of AC drive, the voltage applied to the element is divided to ZnS layer and dielectric layer. Since EL element is structurally equivalent to a series connection of two capacitors, there holds the relation of  $\varepsilon_i V_i/t_i = \varepsilon_z V_z/t_z$  ( $\varepsilon$ : dielectric constant;
- 10 V: voltage applied; t: thickness; suffix i: indicating dielectric; suffix-z: indicating-ZnS), and thus each divided voltage is reversely proportional to the dielectric constant if  $t_i = t_z$ . In said dielectrics such as  $Y_2O_3$ ,  $\varepsilon_i$  is about 4 to 25 and  $\varepsilon_z$  of ZnS is about 9, so
  - that only about 30 to 70% of the whole applied voltage is given to the ZnS layer. In such elements, therefore, a high voltage above 200 V must be applied by a pulse drive of several kHz. Such high voltage gives a great deal of load to the drive circuit and necessitates a
  - specific high-voltage withstanding drive IC, which leads to the increased production cost of the element.

A discussion is here made on what characteristics the dielectric layer is required to have for reducing the drive voltage. From the above-shown relation concerning voltage division, it is noted that the  $\varepsilon_i$  to  $t_i$  ratio  $(\varepsilon_i/t_i)$  must be great. After the start of emission of light, any increment of applied voltage is given to the dielectric layer, so that  $V_{ib}$  (dielectric breakdown voltage

of the dielectric layer) must be also high for giving an excellent dielectric film. Therefore, the figure of merit  $\gamma$  of the dielectric layer is defined as follows:

$$\gamma = \epsilon_i V_{ib} / t_i = \epsilon_i E_{ib}$$

(E<sub>ib</sub>: dielectric breakdown field strength of the dielectric film)

As noted from the above equation,  $\gamma$  is propor-

- 5 tional to the electric charge accumulated per unit area of the dielectric layer when dielectric breakdown occurs. The greater the value of  $\gamma$ , the more stably can be conducted the low-voltage drive. This can be attributed to the following fact. In two EL elements which are same in phosphor layer thickness and dielectric layer thickness but different in properties of dielectric layer (for example, the dielectric layer in one of the elements having the properties of  $\varepsilon_1 = 100$ ,  $E_{\rm ib} = 1 \times 10^6$  V/cm and  $\gamma = 100 \times 10^6$  V/cm while the dielectric layer in another
- element having the properties of  $\epsilon_i$  = 50,  $E_{ib}$  = 3 x 10<sup>6</sup> V/cm and  $\gamma$  = 150 x 10<sup>6</sup> V/cm), naturally the former element can start to emit at a lower voltage than the latter element as they are same in thickness of dielectric layer. However, in the latter element where  $\epsilon_i$  = 50 and  $E_{ib}$  =
- $3 \times 10^6$  V/cm, if it is equalized to the former element in breakdown strength, its layer thickness can be reduced to 1/3. Consequently, its dielectric capacity is

- trebled, boosting  $\epsilon_1$  to 150. Therefore, a greater figure of merit allows the production of an element which emits light at a lower voltage, regardless of  $\epsilon_1$ . The greater the value of  $\gamma$ , the better, but practically,
- it is desirable that  $\gamma$  is about 10 times the value of 14 x 10  $^6$  V/cm that is obtained by substituting  $\varepsilon_z$  = 9 and  $E_{zb}$  = 1.6 x 10  $^6$  V/cm of ZnS for  $\varepsilon_i$  and  $E_{ib}$  in the above-shown formula and used as a standard value for low-voltage luminescence.
- The conventional dielectric films are small in their figure of merit, which is about 50 x  $10^6$  V/cm in the case of  $Y_2O_3$ , about 30 x  $10^6$  V/cm in the case of  $Al_2O_3$  and about 70 x  $10^6$  V/cm in the case of  $Si_3N_4$ , and thus they are not suited for low-voltage luminescence.
- Recently, proposals have been made on use of a thin film mainly composed of  $PbTiO_3$ ,  $Pb(Ti_{1-x}Zr_x)O_3$  or like substance having a high dielectric constant as dielectric layer in an electroluminescent element. These substances are high in  $\varepsilon_i$  which is over 150, but they are
- 20 low in E<sub>ib</sub> which is on the order of 5 x 10<sup>5</sup> V/cm, so that when using these substances, it is required to greatly increase the film thickness in comparison with the conventional dielectric materials. For guaranteeing the reliability of the element produced, it is required that
- the dielectric film has a thickness greater than 15,000 Å, for 6,000 Å in thickness of ZnS film. Generally, in case of using said substances, the grains in the film tend to grow to cause clouding of the film because of large

- 1 film thickness and high substrate temperature at the time
   of formation of the film. In an X-Y matric display using
   such clouded film, light is emitted even from the non luminescent segments as the light from the other
- 5 segments is scattered, resulting in a degraded image quality.

The present inventors had already proposed an EL element using a dielectric film chiefly composed of  $\overline{\text{SrTiO}_3}$ , which dielectric film is high in both  $\overline{\text{E}_{ib}}$  and

- the product of  $E_{ib}$  and  $\varepsilon_i$ , proof against clouding and suited for low-voltage drive. For instance, there had been obtained an  $SrTiO_3$  dielectric film in which  $\varepsilon_i$  = 140 and  $E_{ib}$  = 1.5 MV/cm, the product thereof being greater than that of a BaTiO<sub>3</sub> film (10  $\leq \varepsilon_i \leq$  40,  $E_{ib}$  up to 2 MV/cm).
  - Reduction of driving voltage is desirable for the betterment of reliability and production cost of the drive circuits, but no enough technical breakthrough has been attained in this regard. In order to increase the luminance of the phosphor layer, this layer is subjected to a
  - heat treatment after formation of the film, but in case a dielectric layer is present beneath said phosphor layer, the dielectric layer also undergoes the heat treatment. Consequently, if the dielectric layer thickness is greater than about 0.5  $\mu m$ , certain fault is found to take
  - place in the dielectric film, affecting the breakdown strength of the element. Also, the mode of dielectric breakdown tends to become propagating and is unable to self-heal.

- The present invention is intended to obtain a dielectric film which is better suited for low-voltage drive and also has higher reliability than said SrTiO<sub>3</sub> dielectric film. It is especially envisaged in this
- invention to obtain a dielectric film of the type whose dielectric breakdown, if any, is restricted to self heal, keeping free of propagating breakdown which can be a fatal defect for an EL element.

#### BRIEF DESCRIPTION OF THE DRAWING

The drawing is a schematic sectional view of a thin-film electroluminescent element in an embodiment of this invention.

In the drawing, numeral 1 designates a glass substrate, 2 a transparent electrode, 3 a dielectric film, 4 a ZnS-Mn phosphor film, 5 a Ta<sub>2</sub>O<sub>5</sub> film, 6 a PbNb<sub>2</sub>O<sub>6</sub> film, and 7 an Al electrode.

#### BEST MODE FOR CARRYING OUT THE INVENTION

electroluminescent element comprising a filmy phosphor
layer, a filmy dielectric layer provided on at least one
side of said phosphor layer, and two electrode layers at
least one of which is pervious to light, said electrode
layers being so arranged as to apply a voltage to said
phosphor and dielectric layers, wherein said dielectric
layer is essentially of a composition represented by the
formula:  $x(Ti_{1-y}^{A}v^{O}2) - (1-x)(Ba_{1-z}^{B}z^{O})$  wherein

1 0.4  $\leq$  x  $\leq$  0.8, 0 < y < 1, 0  $\leq$  z < 1, A is at least one element selected from Zr, Hf and Sn, and B is at least one element selected from Mg and Ca.

The present-invention features a novel composition of dielectric film used in the conventional thin-film luminescent elements. According to the present invention, a dielectric film having  $\varepsilon_{i}$  above 50 and  $E_{ib}$  of 3 x 10  $^{6}$  V/cm could be obtained by substituting the position of

Ti in a TiO2-BaO system with Zr, Hf or Sn and further

- above. The film was formed by magnetron RF sputtering method used the sintered ceramic targets prepared for the respective compositions. The result of chemical analysis of the formed film showed the substantial agreement of its composition with that of the target.
  - The dielectric film of said composition and structure has the excellent properties for use in an EL element in comparison with the conventional dielectric films. For instance, in the case of a BaO-TiO $_2$ -SnO $_2$
  - system, the produced film shows higher  $\varepsilon_i$  and  $E_{ib}$  than the conventional BaTiO $_3$  or SrTiO $_3$  film, and accordingly the value of  $\varepsilon_i$  x  $E_{ib}$  is greater than those in said conventional films. Further, the film according to this invention shows no trace of clouding due to the growth of grains and is transparent, so that when it is used as the dielectric layer in an EL element, there can be obtained an EL element with excellent image quality.

- It was also found that the substitution of Ti with Zr or Hf can provide as high  $\varepsilon_i$  or  $E_{ib}$  as in the case of substitution with Sn and enables obtainment of a characteristic heat-resistant dielectric film. Cracking of the
- film in the process of heat treatment leads to a reduction of reliability of the produced EL element because such cracking could cause disconnection of the matrix electrode though such is very rare. Therefore, use of a multiple
  component dielectric film shown here enables the high-
- 10 yield production of high-reliability EL elements free of cracks in the dielectric layer.

The present invention will be further described below by way of the embodiments thereof with reference to the accompanying drawing.

- As illustrated in the drawing, on a glass substrate 1 provided with an ITO transparent electrode 2, a dielectric film 3 having a composition of  $x(\text{Ti}_{0.8}\text{Sn}_{0.2}\text{O}_2) (1-x)\text{BaO was deposited to a thickness of 5,000 Å by magnetron RF sputtering. The sput-$
- tering of said composition was made by changing the value of x: 0.4, 0.5, 0.6, 0.7 and 0.8. A mixed gas of  $O_2$  and Ar (partial pressure of  $O_2$ : 25%) was used as sputtering gas, the gas pressure during sputtering being 0.8 Pa. Used as target was a ceramic plate prepared by mixing
- ingredient powders in said composition and sintering the mixture at 1,400°C. The substrate temperature was 400°C. The produced films with the respective compositions (differing only in the value of x in the above-shown

- l composition) were all transparent and showed no cloudiness. At the point where the dielectric film 3 was formed, the values of  $\epsilon_i$  and  $\epsilon_{ib}$  of the film of each composition were checked. Then ZnS and Mn were simultaneously deposited
- on the dielectric film by electron-beam deposition to form a ZnS-Mn phosphor layer 4 with a thickness of 5,000 Å, and this layer was subjected to a heat treatment in vacuo at 600°C for one hour. For protection of said ZnS-Mn layer, a 400 Å thick Ta<sub>2</sub>O<sub>5</sub> film 5 was further formed on
- 10 said ZnS-Mn layer by electron-beam deposition.—On said  ${\rm Ta_2O_5}$  film 5 was additionally deposited a PbNb $_2{\rm O_6}$  film 6 to a thickness of 1,000 Å by magnetron RF sputtering. An Ar mixed gas containing 25% of  ${\rm O_2}$  was used as sputtering gas. The sputtering gas pressure was 3 Pa. A ceramic of
  - PbNb $_2$ O $_6$  was used as target and the substrate temperature was controlled to 380°C. Finally, a 1,000 Å thick Al film 7 was formed as top electrode by electric resistance heating deposition to complete an EL element.

-Each of the thus formed EL elements was driven

20 by an AC pulse with a repetitive frequency of 5 kHz to determine the voltage-luminance characteristic. Table I shows the electrical properties and luminous characteristics of the elements with the respective dielectric compositions (differing in value of x).

Table 1

No.	x	ε i	E <sub>ib</sub> (V/cm)	ε <sub>i</sub> × Ε <sub>ib</sub> (V/cm)	Lumi- /applied nance voltage (Cd/m <sup>2</sup> ) (V)
1	0.4	60	3.0 x 10 <sup>6</sup>	180 x 10 <sup>6</sup>	3500 / 120
2	0.5	70	3.0 x 10 <sup>6</sup>	210 x 10 <sup>6</sup>	3500 / 110
3	0.6	55	3.2 x 10 <sup>6</sup>	176 x 10 <sup>6</sup>	3500 / 125
4	0.7	40	3.5 x 10 <sup>6</sup>	140 x 10 <sup>6</sup>	3450 / 135
5	0.8	35	3.5 x 10 <sup>6</sup>	112 x 10 <sup>6</sup>	3400 / 140

The voltage at which the saturation brightness of 3,400 to 3,500 Cd/m² is reached is given in the table as a measure of luminous characteristics. As seen from the table, the dielectric constant is maximized and also the value of  $\epsilon_i$  x  $E_{ib}$  becomes largest when x is 0.5. What is especially noteworthy in this system is that the dielectric breakdown field strength  $E_{ib}$  is above 3 x  $10^6$  V/cm, which is far greater than that in the case of SrTiO3, and that the mode of dielectric breakdown is

showed a dielectric constant above 100 when the heat treatment after deposition was conducted at 600°C for one hour. From the above-shown dependency of  $\varepsilon_i$  and  $E_{ib}$  on the compositional change (in x), it was found that the produced films are far higher in  $E_{ib}$  than that of SrTiO $_3$  and has a substantially same figure of merit as that of SrTiO $_3$  when x is between 0.4 and 0.8. Being high in  $E_{ib}$  is an essential factor for improving the

self-healing type. Also, some of the obtained elements

10

- 1 reliability of the thin-film electroluminescent element. As regards the luminous characteristics, the voltage that provides the saturation brightness of 3,400-3,500  $Cd/m^2$  is minimized (110 V) when x = 0.5, indicating low-
- 5 voltage drivability of the element. Also, said voltage is below 140 V even when x is in the range of 0.4 to 0.8, and thus low-voltage drive is possible.

Considering the above-shown results synthetical-

10 by the formula:  $x(Ti_{0.8}Sn_{0.2}O_2) - (1-x)BaO$ , with x in the formula being defined as  $0.4 \le x \le 0.8$ , it is possible to obtain a film more excellent than the conventional  $SrTiO_3$  or  $BaTiO_3$  film as a dielectric film for a low-voltage drive type EL element.

- We will now describe the case where the amount of partial substitution of Ti with Sn is further changed in said  $x(Ti_{0.8}Sn_{0.2}O_2) (1-x)BaO$  system by defining x to 0.5 at which the most excellent film properties are obtained. The amount of substitution with Sn was varied
- from 0 to 0.4. The method of evaluation of dielectric film, the construction and preparing conditions of the element and the conditions for determination of luminous characteristics are the same as in the case of said

  Ti<sub>0.8</sub>Sn<sub>0.2</sub>O<sub>2</sub> BaO system. Table 2 shows the results obtained when substituting Sn for the position of Ti.

  In the table, there was given a new item indicating the percentage of cracking of the dielectric film at the time

of annealing of the ZnS-Mn phosphor layer 4 formed on

- the dielectric film 3 (the percentage of cracking was calculated from the number of samples which were cracked in 10 pieces of samples tested). There was also given a column for qualitatively showing whether the state-of
- dielectric breakdown is self-healed or not by observing the mode of dielectric breakdown after determination of dielectric breakdown field strength.

Table 2

No.	У	ε i	E <sub>ib</sub> (V/cm)	ε <sub>i</sub> x E <sub>ib</sub> (V/cm)	Parcen- tage of cracked samples (%)	Dielectric breakdown was: Self- healed (o), Not self- healed (x)
6	0	63	1.5 x 10 <sup>6</sup>	95 x 10 <sup>6</sup>	100	x
7	0.1		1.6 x 10 <sup>6</sup>		0	0
8	0.2	70	$3.3 \times 10^6$			0
9	0.3	48	3 x 10 <sup>6</sup>		1	0
10	0.4	30	3 x 10 <sup>6</sup>	90 x 10 <sup>6</sup>	0	0

As seen from Table 2, both  $\epsilon_{\dot{1}}$  and  $E_{\dot{1}\dot{b}}$  tend to increase with the partial substitution of Ti with Sn.

Thus, when Ti is substituted with Sn to a degree where y is 0.3 or below, there can be obtained a greater figure of merit than that obtainable with a  $0.5\text{TiO}_2$  - 0.5BaO film. Especially, the figure of merit of  $\epsilon_i$  x  $\epsilon_{ib}$  is maximized when the substitution rate y of Sn is 0.2 or

- the high-yield production of low-voltage drive EL element proof against cracking at the time of annealing is possible. Also, when y (Sn substitution rate) was
- o.1, 0.2 and 0.3, the dielectric constant of the dielectric film after annealing was 150, 130 and 100, respectively, indicating a further reduction of drive voltage for EL element by the Sn substitution for Ti in said range.
- The effect of similar substitution for Ti with Zr and Hf was examined in the same way as in the case of substitution with Sn. It was found that, in this case, the value of  $\epsilon_i$  x  $E_{ib}$  is maximized and also the film becomes most resistant to cracking when y is 0.2 or
- therearound as in the case of Sn. It is especially noteworthy in the case of Zr and Hf that the range of substitution rate (y) that provides a large figure of merit is wide, and it was confirmed that even when y was 0.5, the produced dielectric film could well serve
- for a low-voltage drive EL element. For instance, in the case of  $0.5(\text{Ti}_{0.7}\text{Zr}_{0.3}\text{O}_2)-0.5\text{BaO}$ ,  $\varepsilon_i$  was 60 and  $\varepsilon_{ib}$  was 2.5 x  $10^6$  V/cm, and further the mode of dielectric breakdown was self-healing type. Also, in the case of  $0.5(\text{Ti}_{0.5}\text{Zr}_{0.5}\text{O}_2)-0.5\text{BaO}$ ,  $\varepsilon_i=30$  and  $\varepsilon_{ib}=3\times10^6$  V/cm, and in the case of  $0.5(\text{Ti}_{0.6}\text{Hf}_{0.5}\text{O}_2)-0.5\text{BaO}$ ,

It is needless to say that an excellent low-voltage drive EL element can be obtained by properly

 $\varepsilon_i = 35$  and  $\varepsilon_{ib} = 3 \times 10^6$  V/cm.

l combining the substituting elements Sn, Zr and Hf for the position of Ti.

The results obtained from substitution for the position of Ba with Mg and Ca are shown below.

- The method of evaluation of dielectric film, the structure and preparing conditions of the element and the luminous characteristic determining conditions were the same as in the case of said  $0.5 \text{Ti}_{1-y} \text{Sn}_{y}^{0} \text{O}_{2} 0.5 \text{BaO}$  system.
- Table 3 shows the results obtained from Mg substitution for the position of Ba.

5

Table 3

No.	Valve of z in 0.5 Ti <sub>0.8</sub> Sn <sub>0.1</sub> O <sub>2</sub> - 0.5(Ba <sub>1-z</sub> Mg <sub>z</sub> O)	Ēi		ε <sub>i</sub> x E <sub>ib</sub> (V/cm)	Parcen- tage of crack- ing
11	,		1.6 x 10 <sup>6</sup>		0
12	0.1	75	i .	135 x 10 <sup>6</sup>	0
13	-0.2	<del>- 70</del>	$2.0 \times 10^{6}$		4
14	0.3	65	$2.5 \times 10^6$		1
15	0.4	50	$2.8 \times 10^6$		l .
16	0.6	: 20	$3.0 \times 10^6$	60 x 10 <sup>6</sup>	0
		1	<u> </u>		

As seen from Table 3, partial substitution of Ba with Mg produces an decreasing tendency of  $\epsilon_i$  while causing an increase of  $E_{ib}$ . Thus, in the range of about 10 to 30% substitution for Ba with Mg, the produced film

- is improved in figure of merit over the non-substituted film. Also, no cracking was caused by the substitution of Ba with Mg. However, when the Mg substitution rate exceeds 60%, the dielectric constant is reduced to the
- order of 20 and the figure of merit falls below the level of 100 x 10<sup>6</sup> V/cm suited for low-voltage luminescence (about 7 times the figure of merit provided by ZnS).

  Therefore, the appropriate substitution rate of Mg for

  Ba would be less than 40%. In this range, it is
- 10 possible to produce, in a high yield, a low-voltage drive EL element having no risk of cracking at the time of annealing.

The case of Ca substitution in the completely same manner as in the case of Mg described above was also examined. In this case, there was observed the same tendency concerning  $\epsilon_i$  and  $E_{ib}$  as in the case of Mg, and also no crack was seen in the produced film. The optimal range of Ca substitution rate was determined to be less than 30%; any greater substitution rate drops the figure

of merit below 100 x  $10^6$  V/cm and also makes the film prone to clouding. In the film with a composition of  $0.5(\text{Ti}_{0.9}\text{Sn}_{0.1}^{0}) - 0.5(\text{Ba}_{0.7}^{\text{Ca}_{0.3}^{0}})$ ,  $\epsilon_{i} = 60$ ,  $\epsilon_{ib} = 2.3 \times 10^6$  V/cm, and  $\epsilon_{i} \times \epsilon_{ib} = 138 \times 10^6$  V/cm. It was also found that the produced dielectric film comes to show a self-healing type dielectric breakdown when the position of Ti is substituted with Sn by a factor of about 0.1 to 0.3.

As described above, by use of the compositions

l according to this invention, there can be obtained a dielectric film which is proof against cracking and is characteristically high in s<sub>i</sub> and E<sub>ib</sub> and hence also high in figure of merit. Further, when Ti in the composition

5 tion is substituted with Sn, Zr or Hf, dielectric breakdown of the film is rendered self-healing type.

It is of course possible to adopt a fourcomponent system incorporating said substitutions within
the specified range of substitution rate for the purpose

10 of combining the advantages of the respective substituting
elements (Sn, Zr or Hf for Ti, and Mg or Ca for Ba) in
the basis composition of TiO<sub>2</sub>-BaO.

### INDUSTRIAL APPLICABILITY

As described above, according to the present invention, the filmy dielectric layer of a thin-film electroluminescent element is composed of a dielectric having a composition of  $x(Ti_{1-y}^Ay^O_2) - (1-x)BaO$  which is high in figure of merit and resistant to cracking and whose dielectric breakdown tends to self-heal, so that it is possible to obtain a low-voltage drive type electroluminescent element with high image quality and reliability in a high yield. This is of great industrial value from the aspects of improvement of reliability and production cost of drive circuits.

#### WHAT IS CLAIMED IS:

a filmy phosphor layer, a filmy dielectric layer provided on at least one side of said phosphor layer, and two electrode layers at least one of which is pervious to light, said electrode layers being so arranged as to apply a voltage to said phosphor and dielectric layers, wherein the essential composition of said dielectric layer is expressed by the following formula:

# $-x(Ti_{1-y}A_yO_2)$ ---(1-x) $(Ba_{1-z}B_zO)$

wherein x, y and z are the numbers defined as:  $0.4 \le x \le 0.8$ , 0 < y < 1, and  $0 \le z < 1$ , and A is at least one element selected from Zr, Hf and Sn, and B is at least one element selected from Mg and Ca.

- 2. A thin-film electroluminescent element according to Claim 1, wherein in the compositional formula of the essential components of the dielectric layer, A is Zr or Hf and y is a range defined as: y < 0.5.
- 3. A thin-film electroluminescent element according to Claim 1, wherein in the compositional formula of the essential components of the dielectric layer, A is Sn and y is a range defined as:  $y \le 0.3$ .
- 4. A thin-film electroluminescent element according to Claim 1, wherein in the compositional formula of the essential components of the dielectric layer, B is Mg and z is a range defined as:  $z \le 0.4$ .

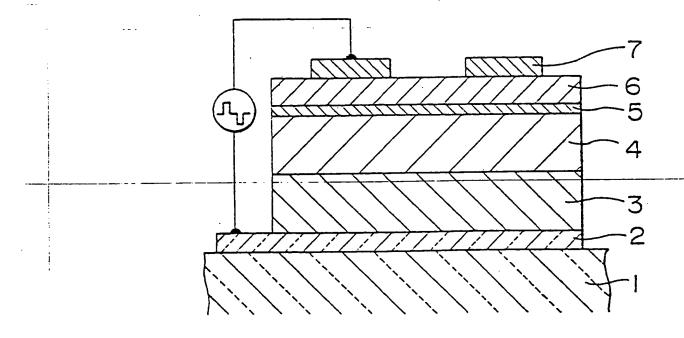
5. A thin-film electroluminescent element according to Claim 1, wherein in the compositional formula of the essential components of the dielectric layer, B is Ca and z is a range defined as:  $z \le 0.3$ .

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## EUROPEAN SEARCH REPORT

EP 84 30 8539

ategory	Citation of document with in of relevant	ERED TO BE RELEVAN  dication, where appropriate, passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int. CI.4)
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Y	FR-A-2 376 500 ( * Claims 1-6 *	SIEMENS)	1	
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